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54 **Rhodium-catalyzed olefin dimerization.**

57 A process is disclosed for preparing functionalized linear olefins by dimerizing terminal olefins in the presence of a cationic rhodium compound. Novel rhodium compounds useful in this process are also disclosed.

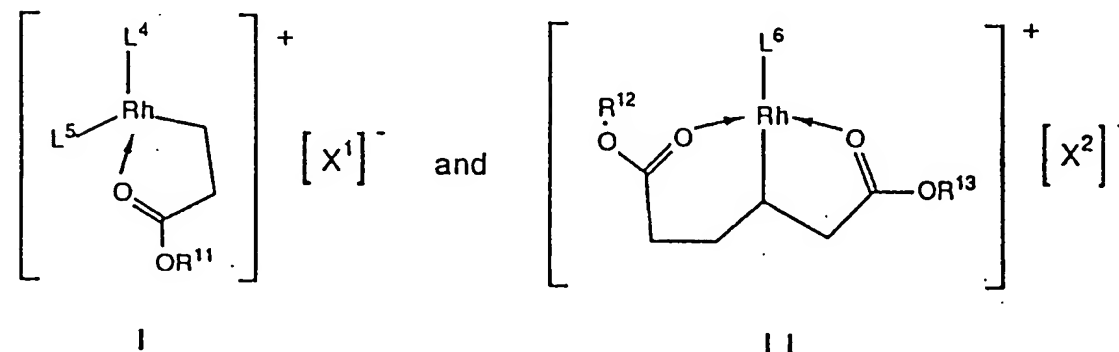
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5 BACKGROUND OF THE INVENTION

Brookhart et al. (J. Amer. Chem. Soc. **1988**, *110*, 8719-8720) disclose the use of a cationic rhodium catalyst containing a pentamethylcyclopentadienyl ligand in the dimerization of ethylene to butenes.

SUMMARY OF THE INVENTION

This invention also provides novel compounds, I and II, which are useful in the process of this invention,



L⁴ is an anionic pentahapto ligand;
L⁵ is a neutral 2-electron donor ligand;

- R^{11} is selected from the group of C_1 - C_{10} alkyl;
 $[X^1]^-$ is a non-coordinating anion;
 L^6 is an anionic pentahapto ligand;
 R^{12} and R^{13} are independently selected from the group consisting of C_1 - C_{10} alkyl; and
 $[X^2]^-$ is a non-coordinating anion.

DETAILED DESCRIPTION OF THE INVENTION

The process of this invention can be used to homodimerize or codimerize functionalized terminal olefins in a linear, tail-to-tail fashion, or to dimerize functionalized terminal olefins with terminal alkenes. The products of the process of this invention are linear, functionalized olefins in which a carbon-carbon bond has been formed between the methylene carbons of the olefin reactants. Specific examples of useful products include dialkyl hexenedioates, which are precursors to adipic acid.

In the process of this invention, a linear functionalized olefin is prepared by reacting a first terminal olefin, $CH_2=CR^1R^2$, with a second terminal olefin, $CH_2=CR^3R^4$, in the presence of a cationic rhodium compound, $[L^1RhL^2L^3R]^+X^-$; wherein

- R^1 is selected from the group consisting of H and C_1 - C_{10} alkyl;
 R^2 is selected from the group consisting of H, C_1 - C_{10} alkyl, phenyl, C_7 - C_{12} alkyl-substituted phenyl, $-COOR^5$, $-C(O)NR^6R^7$, and $-C(O)H$;
 R^3 is selected from the group consisting of H and C_1 - C_{10} alkyl;
 R^4 is selected from the group consisting of $-COOR^8$, $-C(O)NR^9R^{10}$, and $-C(O)H$;
 R^5 and R^8 are independently selected from the group consisting of C_1 - C_{10} alkyl;
 R^5 , R^7 , R^9 , and R^{10} are independently selected from the group consisting of H and C_1 - C_{10} alkyl;
 L^1 is an anionic pentahapto ligand;
 L^2 and L^3 are neutral 2-electron donor ligands;
 R is selected from the group of H, C_1 - C_{10} alkyl, C_6 - C_{10} aryl, and C_7 - C_{10} aralkyl ligands;
 X^- is a non-coordinating anion; and
 wherein two or three of L^2 , L^3 and R are optionally connected.

Suitable terminal olefins, $H_2C=CR^1R^2$, include: ethylene; terminal alkenes containing 3-12 carbon atoms, e.g., propene, 1-butene, isoprene, 1-pentene, 1-hexene, and 1-heptene; styrene; 4-methylstyrene; alkyl acrylates, where the alkyl group contains 1-10 carbon atoms, e.g., methyl acrylate and ethyl acrylate; methyl methacrylate; acrylamide; methacrylamide; N-alkyl acrylamides, where the alkyl group contains 1-10 carbon atoms, e.g., N-methylacrylamide; N-methyl methacrylamide; N,N-dialkyl acrylamides, where the alkyl groups contain 1-10 carbon atoms, e.g., N,N-dimethylacrylamide; acrolein; and methacrolein.

Suitable functionalized terminal olefins, $H_2C=CR^3R^4$, include: alkyl acrylates, where the alkyl group contains 1-10 carbon atoms, e.g., methyl acrylate and ethyl acrylate; methyl methacrylate; acrylamide; methacrylamide; N-alkyl acrylamides, where the alkyl group contains 1-10 carbon atoms, e.g., N-methylacrylamide; N-methyl methacrylamide; N,N-dialkyl acrylamides, where the alkyl groups contain 1-10 carbon atoms, e.g., N,N-dimethylacrylamide; acrolein; and methacrolein.

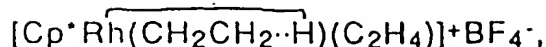
Preferably, $H_2C=CR^1R^2$ is ethylene, propylene, styrene, methyl acrylate, ethyl acrylate, acrolein, or N,N-dimethyl acrylamide. Preferably, $H_2C=CR^3R^4$ is methyl acrylate, ethyl acrylate, acrolein, or N,N-dimethyl acrylamide. More preferably, $H_2C=CR^1R^2$ is ethylene, styrene, methyl acrylate or ethyl acrylate and $H_2C=CR^3R^4$ is methyl acrylate or ethyl acrylate. Most preferably, $H_2C=CR^1R^2$ and $H_2C=CR^3R^4$ are both either methyl acrylate or ethyl acrylate.

The terminal olefins, $H_2C=CR^1R^2$ and $H_2C=CR^3R^4$, can be chosen to be the same or different olefins to give, respectively, homodimers or codimers. The efficiency of the production of codimers may depend on the specific olefins chosen, and thus it may be necessary to use a large excess of one of the olefins to obtain the desired codimer.

The cationic rhodium compound used in the process of this invention can be formed in one of several ways. A particularly convenient route involves reacting a precursor, $L^1RhL^2L^3$, with an acid, H^+X^- , where

- L^1 is an anionic pentahapto ligand;
 L^2 and L^3 are neutral, 2-electron donor ligands, or L^2 and L^3 are connected to form a neutral, 4-electron ligand; and
 X^- is a non-coordinating anion.

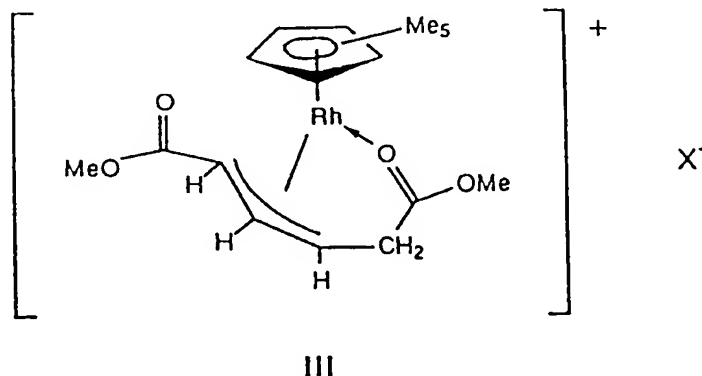
For example, $Cp^*Rh(C_2H_4)_2$ reacts with HBf_4 to give



which is useful in the process of this invention. (Cp^* is pentamethylcyclopentadienyl.) Similarly, compound
 5 **1a** (L^4 is Cp^* ; L^5 is $\text{P}(\text{OMe})_3$; R^{11} is Me; and $[\text{X}']^-$ is BF_4^-) can be made by reacting HBF_4 with $\text{Cp}^*\text{Rh}(\text{P}(\text{OMe})_3)(\text{CH}_2=\text{CHCO}_2\text{Me})$. In these routes to cationic rhodium compounds, suitable acids, H^+X^- , include: HBF_4 ; HPF_6 ; H_2SO_4 ; $\text{CF}_3\text{SO}_3\text{H}$; $\text{CF}_3\text{CO}_2\text{H}$; and tetraarylboronic acids, e.g., HBPh_4 and $\text{HB}(3,5\text{-bis}(\text{trifluoromethyl})\text{phenyl})_4$.

Alternatively, $\text{L}^1\text{RhL}^{2'}(\text{R})\text{Y}$, where Y is a halide and L^1 , $\text{L}^{2'}$, and R are as defined above, can be reacted
 10 with a Lewis acid in the presence of an olefin to form a cationic rhodium compound which is useful in the process of this invention. For example, $\text{Cp}^*\text{Rh}(\text{P}(\text{OMe})_3)(\text{Me})\text{Br}$ could be reacted with AgBF_4 in the presence of methyl acrylate to give the desired cationic rhodium compound, $[\text{Cp}^*\text{Rh}(\text{P}(\text{OMe})_3)(\text{CH}_2=\text{CHCO}_2\text{Me})(\text{Me})]^+\text{BF}_4^-$. In catalyst preparations of this type, suitable Lewis acids include: Ag^+X^- , AlX''_3 , BX''_3 , FeX''_3 , and SbX''_5 , where X'' is halide.

In a third general route, precursors such as $[\text{L}^1\text{RhL}^{2'}\text{L}^4]^+$, where L^4 is a π -allylic ligand and L^1 and $\text{L}^{2'}$
 15 are as defined above, can be reacted with H_2 to give cationic rhodium compounds which are useful in the process of this invention. For example, compounds of the class $[\text{Cp}^*\text{Rh}(\text{MeOC}(\text{O})\text{CH}_2\text{CHCHCHCO}_2\text{Me})]^+\text{X}^-$
 III,



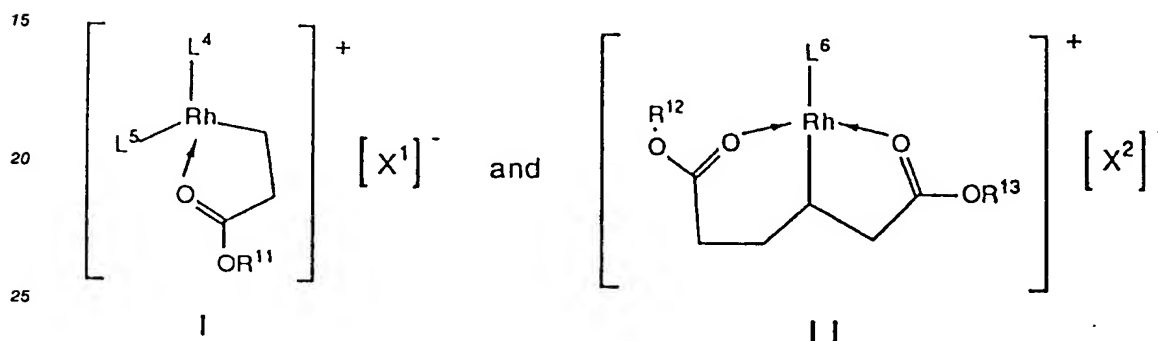
can be reacted with hydrogen to give cationic rhodium compounds which are useful in the process of this
 35 invention. A particularly useful precursor of this type is $[\text{Cp}^*\text{Rh}(\text{MeOC}(\text{O})\text{CH}_2\text{CHCHCHCO}_2\text{Me})]^+[\text{B}(3,5\text{-bis}(\text{trifluoromethyl})\text{phenyl})_4]^-$ IIIa.

In all of these rhodium compounds, suitable pentahapto ligands, L^1 , L^4 and L^5 include: cyclopentadienyl
 and substituted derivatives of cyclopentadienyl containing 1-5 substituents chosen from $\text{C}_1\text{-C}_4$ alkyl,
 40 trifluoromethyl, $\text{C}_6\text{-C}_{10}$ aryl, COOR^{14} (where R^{14} is $\text{C}_1\text{-C}_4$ alkyl), and $\text{C}(\text{O})\text{R}^{15}$ (where R^{15} is $\text{C}_1\text{-C}_4$ alkyl);
 indenyl; fluorenyl; and carboranyl ligands such as (7,8,9,10,11- η)undecahydro-7,8-dicarbaundecaborato(2-)
 and (7,8,9,10,11- η)undecahydro-7,9-dicarbaundecaborato(2-). Preferably, L^1 , L^4 and L^5 are alkyl-substituted
 derivatives of cyclopentadienyl; most preferably, L^1 , L^4 and L^5 are pentamethylcyclopentadienyl (Cp^*).

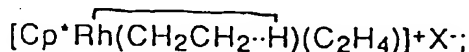
Suitable neutral, 2-electron donors, L^2 , L^3 , $\text{L}^{2'}$, $\text{L}^{3'}$, and L^5 include: carbon monoxide; alkyl-, aryl-, or
 45 mixed alkyl,arylphosphines (e.g., trimethylphosphine, triphenylphosphine, or diethylphenylphosphine); alkyl-,
 aryl-, or mixed alkyl,arylphosphites (e.g., trimethylphosphite, triphenylphosphite, or dimethylphenyl-
 phosphite); olefins (e.g., ethylene, propylene, 1-hexene, 1-octene, methyl acrylate, ethyl acrylate, or
 dimethyl hexenedioate); nitriles (e.g., acetonitrile or benzonitrile); and the carbonyl groups of ketones (e.g.,
 50 acetone) and esters (e.g., methyl acrylate). L^2 and L^3 can be the same or different, provided that if L^2 is a
 phosphine or phosphite, then L^3 is not a phosphine or phosphite. Similarly, $\text{L}^{2'}$ and $\text{L}^{3'}$ can be the same or
 different, but cannot both be phosphine or phosphite ligands. Preferred 2-electron donors include carbon
 monoxide, ethylene, trimethylphosphite, methylacrylate and dimethyl hexenedioate.

Alternatively, L^2 and L^3 , or $\text{L}^{2'}$ and $\text{L}^{3'}$, may be connected to form a neutral 4-electron donor ligand
 which contains two 2-electron-donor sites (olefin, phosphine, phosphite, nitrile or carbonyl groups). Suitable
 55 4-electron-donor ligands of this type include: butadiene, 1,5-pentadiene, methyl vinyl ketone and ac-
 rylonitrile. Similarly, R and L^2 (or $\text{L}^{2'}$) can be connected, as in

The novel compounds, I and II,



Other preferred cationic rhodium compounds include:



Suitable temperatures for the process of this invention range from about -100°C to about 150°C .

depending on the specific catalyst, olefin(s) and pressure. More preferably, the temperature is between 0 °C and 100 °C; most preferably between 20 °C and 80 °C.

The process of this invention is not particularly sensitive to pressure, and pressures of 0.1 atm to 1,000 atm are suitable.

The process of this invention can be conducted in the presence of inert gases such as nitrogen, argon, helium, carbon dioxide and saturated hydrocarbons such as methane. In the preferred mode, the process is conducted in the presence of hydrogen, where the partial pressure of hydrogen is from about 0.1 atm to about 10 atm. Surprisingly, high yields of dimers are obtained and less than 3% saturated products are observed even under 1 atm hydrogen.

EXAMPLES

The following examples are provided to illustrate the invention and are not to be construed as limiting the invention. All preparative manipulations were carried out using conventional Schlenk techniques. Methylene chloride was distilled from P₂O₅ under a nitrogen atmosphere. Methyl acrylate was stored under 4 Å molecular sieves. The rhodium complexes were prepared according to published procedures.

Reaction mixtures were analyzed by ¹H NMR spectroscopy. This method is advantageous since the fate of the rhodium species as well as the conversion of methyl acrylate into dimers can be monitored. The only dimers observed in all cases were linear, tail-to-tail dimers which included E- and Z-CH₃OC(O)-CH=CH-CH₂-CH₂-CO₂CH₃ (from here on referred to as E-2 and Z-2) and E-CH₃OC(O)-CH₂-CH=CH-CH₂-CO₂CH₃ - (from here on referred to as E-3). Normally, the E-2 isomer was the major isomer. Small amounts of E-3 often appeared at the end of the reaction, probably due to isomerization of the E-2 and Z-2 isomers under the reaction conditions. The turnover number (TON) was defined as the number of moles of methyl acrylate consumed/mole of rhodium complex. The most efficient reactions were carried out under 1 atm H₂. Under these conditions very little (<3%) hydrogenation of methylacrylate occurs.

Examples 1-3 demonstrate relatively inefficient dimerization employing Cp*Rh(C₂H₄)(P(OMe)₃) as starting material. In all these examples the reaction was followed by ¹H NMR using NMR tubes sealed under vacuum.

Example 1

HBf₄·Me₂O (32 μL, 0.287 mmol) in 5 mL diethyl ether was added at -30 °C to Cp*Rh(C₂H₄)(P(OMe)₃) (84 mg, 0.215 mmol) in 25 mL ether. The rhodium hydride salt [Cp*Rh(C₂H₄)(P(OMe)₃)H]⁺[BF₄]⁻ precipitated immediately. The mixture was cooled to -80 °C and the ether solution was removed via cannula. The solid was washed with 2 portions of 5 mL of cold diethyl ether and dried under vacuum at low temperature.

Methyl acrylate (7.2 μL, 0.08 mmol) was added to an NMR tube at liquid nitrogen temperature containing [Cp*Rh(C₂H₄)(P(OMe)₃)H]⁺[BF₄]⁻ (8 mg, 0.017 mmol) in 0.6 mL CD₂Cl₂. The NMR tube was then sealed under vacuum. The reaction was monitored by ¹H NMR. A new complex Cp*Rh-(CH₂CH₂CO₂Me)(P(OMe)₃) was obtained and slow dimerization of the methyl acrylate was observed. (50% conversion after 9 days)

Example 2

The new complex [Cp*Rh(CH₂CH₂CO₂Me)(P(OMe)₃)]⁺BF₄⁻ (Ia) was prepared starting from [Cp*Rh-(C₂H₄)(P(OMe)₃)H]⁺[BF₄]⁻ (140 mg, 0.293 mmol) and methyl acrylate (36 μL, 0.40 mmol) in 5 mL CH₂Cl₂. Then methyl acrylate (250 μL, 2.78 mmol) was added at room temperature. Slow dimerization was obtained: 17% conversion after 24 h and 58% after 12 days.

NMR data for [Cp*Rh(CH₂CH₂CO₂Me)(P(OMe)₃)]⁺BF₄⁻ (Ia): ¹H NMR (CD₂Cl₂, 400 MHz, 23 °C): δ 3.79 (s, CO₂CH₃), 3.71 (d, J_{P-H} = 12 Hz, P(OCH₃)₃), 2.9 (m, CH₂), 2.2 (m, CH₂), 1.67 (d, J_{P-H} = 4 Hz, C₅(CH₃)₅). ¹³C{¹H} NMR (CD₂Cl₂, 100 MHz, 23 °C): δ 191.0 (s, CO₂CH₃), 101.2 (s, C₅(CH₃)₅), 55.6 (s, CO₂CH₃), 53.2 (d, J_{P-C} = 4 Hz, P(OCH₃)₃), 39.1 (s, CH₂CO₂CH₃), 13.0 (t, J_{Rh-C} = J_{P-C} = 18 Hz, RhCH₂), 9.3 (s, C₅(CH₃)₅).

Example 3

Methyl acrylate (77 μL, 0.86 mmol) was added to [Cp*Rh(C₂H₄)(P(OMe)₃)H]⁺[BF₄]⁻ (12 mg, 0.025 mmol) prepared following the method described in Example 1. After 4 days, 50% conversion to dimers was obtained.

Examples 4-13 demonstrate very efficient dimerization employing Cp*Rh(C₂H₄)₂ as starting material.

Only linear dimers were obtained.

Example 4

5 HBF₄•OMe₂ (one drop) was added at -40 °C to Cp*Rh(C₂H₄)₂ (6 mg, 0.02 mmol) in 0.5 mL of CD₂Cl₂ in an NMR tube. After shaking, the tube was frozen at liquid nitrogen temperature. Methyl acrylate (250 µL, 2.78 mmol) was added and then the tube was sealed under vacuum at liquid nitrogen temperature. The reaction was then followed by NMR analysis at room temperature. After 45 min, 97% conversion to dimers was obtained. Dimers: E-2, 94%; Z-2, 4%; E-3, 2%.

Example 5

15 HBF₄•OMe₂ (one drop) was added at -50 °C to Cp*Rh(C₂H₄)₂ (6 mg, 0.02 mmol) in 5 mL of CH₂Cl₂ in a 100 mL Schlenk flask. Methyl acrylate (2.5 mL, 27.8 mmol) (degassed under N₂) was added at -50 °C. The mixture was then stirred at 0 °C. The reaction was followed by NMR by withdrawing 50 µL of the mixture and adding it to 0.5 mL of CD₂Cl₂. After 20 h at 0 °C, 63% conversion to dimers was obtained. Dimers: E-2, 86%; Z-2, 14%. TON = 876.

Example 6

20 The procedure described in Example 5 was repeated, except that the mixture was kept in a water bath at room temperature. After 3.83 h, 67% conversion to dimers was obtained. Dimers: E-2, 85%; Z-2, 18%. TON = 931.

In Examples 7-11 and 13, HBPh₄** indicates HB[3,5-bis(trifluoromethyl)phenyl]₄.

Example 7

25 HBPh₄**•(Et₂O)₂ (29 mg, 0.029 mmol) was added to Cp*Rh(C₂H₄)₂ (6 mg, 0.020 mmol) in 5 mL CH₂Cl₂ at 0 °C. Methyl acrylate (3 mL, 33.3 mmol) was added at 0 °C and after 5 min the Schlenk flask was kept at room temperature in a water bath. Results are presented in the following table.

Time (h)	%Conversion to dimers
0.25	5
1	16
3	45
6	62
24	75

40 At 24 h, dimers were: E-2, 89%; Z-2, 11%. TON = 1249.

Example 8

45 This example shows that the presence of a solvent is not necessary.

HBPh₄**•(Et₂O)₂ (49 mg, 0.048 mmol) in 2 mL of diethyl ether was added at 0 °C to Cp*Rh(C₂H₄)₂ (10 mg, 0.034 mmol) in 2 mL of diethyl ether. After stirring 7 min, the mixture was evaporated to dryness at 0 °C under vacuum. Then methyl acrylate (8 mL, 88.9 mmol) was added at 0 °C to the remaining solid. After stirring 5 min, the Schlenk flask was kept in a water bath at room temperature. 47% conversion was obtained: E-2, 88%; Z-2, 12%. TON = 1229.

Example 9

55 This example shows that dimerization occurs at a temperature as low as -80 °C.

HBPh₄**•(Et₂O)₂ (38 mg, 0.037 mmol) in 0.3 mL CD₂Cl₂ was added at 0 °C to Cp*Rh(C₂H₄)₂ (7 mg, 0.024 mmol) in 0.5 mL CD₂Cl₂ in an NMR tube. After cooling to -80 °C, methyl acrylate (20 µL, 0.222 mmol) was added, and the tube was shaken just before its introduction into the NMR probe. Dimerization was observed at -80 °C, but the reaction was very slow.

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In Examples 10-13, the reactions were performed using N₂ and H₂ atmospheres.

Example 10

5 HBPh₄^{***}(Et₂O)₂ (49 mg, 0.048 mmol) in 2 mL CH₂Cl₂ was added at 0 °C to Cp*Rh(C₂H₄)₂ (10 mg, 0.034 mmol) in 10 mL CH₂Cl₂. After stirring 10 min, methyl acrylate (8 mL, 88.9 mmol) was added to the mixture. The Schlenk flask was then kept at room temperature in a water bath. After 4 h reaction under N₂ atmosphere, 36% conversion to dimers was obtained. At this point, the mixture was divided into two fractions: one fraction was kept under N₂ and 47% conversion was finally obtained. H₂ was bubbled through
10 the second fraction for 1 h, and 95% conversion to dimers was finally obtained. TON = 2483 (H₂ atmosphere).

Example 11

15 HBPh₄^{***}(Et₂O)₂ (50 mg, 0.049 mmol) in 1.5 mL CH₂Cl₂ was added at 0 °C to Cp*Rh(C₂H₄)₂ (10 mg, 0.034 mmol) in 2.5 mL CH₂Cl₂. After stirring 10 min, methyl acrylate (20 mL, 222.3 mmol) was added to the solution. The Schlenk flask was then kept at room temperature in a water bath under H₂ atmosphere. The results are reported in the following table:

20

Time (h)	%Conversion to dimers
4.33	14
22.33	68
48	>99.9

25

At 48 h, TON = 6538.

Turnover rate = 3.5 mol CH₂=CHCO₂Me/mol(Rh)/min at 25 °C.

Dimers: E-2, 95%; Z-2, 3%; E-3, 2%.

30

Example 12

One drop of HBF₄·Me₂O was added at -40 °C to Cp*Rh(C₂H₄)₂ (10 mg, 0.034 mmol) in 10 mL of CH₂Cl₂. Methyl acrylate (8 mL, 88.9 mmol) was added to the mixture, and the mixture was then heated to 40-50 °C under an H₂ atmosphere. (The Schlenk flask was equipped with a water condenser.) The reaction
35 was only monitored for 4 h and at that point, 69% conversion was obtained.

Turnover rate = 7.5 mol CH₂=CHCO₂Me/mol(Rh)/min at 40 °C.

Example 13

40

HBPh₄^{***}(Et₂O)₂ (50 mg, 0.049 mmol) in 3 mL CH₂Cl₂ was added at 0 °C to Cp*Rh(C₂H₄)₂ (10 mg, 0.034 mmol) in 3 mL CH₂Cl₂. After stirring 10 min, methyl acrylate (20 mL, 222.3 mmol) was added to the solution. The Schlenk flask was then kept at room temperature in a water bath under H₂ atmosphere. The results are reported in the following table:

45

50

55

Time (h)	%Conversion to dimers
2	12
3.25	20
4.33	27
5.33	33
7.75	47
9.75	59
11.50	67
12.92	75
14.83	84
16.75	91
18.50	95
20.33	97

After 11.50 h, the mixture was frozen in dry ice/acetone overnight. Just after thawing, no change was noticed in the monomer/dimer ratio and the reaction was then monitored in the same manner as before freezing. After 36 h at room temperature, >99.9% conversion was obtained, giving a TON = 6538. (No data points were taken between 20.33 and 36 h.)

Turnover rate = 6.6 mol CH₂ = CHCO₂Me/mol(Rh)/min at 25 °C (over the initial 10 h period).

Dimers: E-2, 94%; Z-2, 5%; E-3, 1%.

Example 14

The procedure described in Example 13 was repeated, except that the mixture was heated to 60 °C under H₂ atmosphere.

After 3 h, 94% conversion was obtained. An additional 20 mL of methyl acrylate was added, and after 22 h, 99% conversion was obtained at 60 °C, giving a TON = 13,000.

Turnover rate = 65 mol CH₂ = CHCO₂Me/mol(Rh)/min at 60 °C (over the initial (1 h) period).

Products: Dimers (98%): E-2, 93%; Z-2, 6%; E-3, 1%

Methyl propionate (2%).

Example 15

This example describes the synthesis (2 methods) and the characterization of the new complexes [Cp*RhCH(CH₂CO₂Me)(CH₂CH₂CO₂Me)]⁺[BPh₄⁻]⁻ (IIb) and {Cp*Rh(η³-MeOC(O)CH₂CHCHCHCO₂Me)}⁺[BPh₄⁻]⁻ (IIIb).

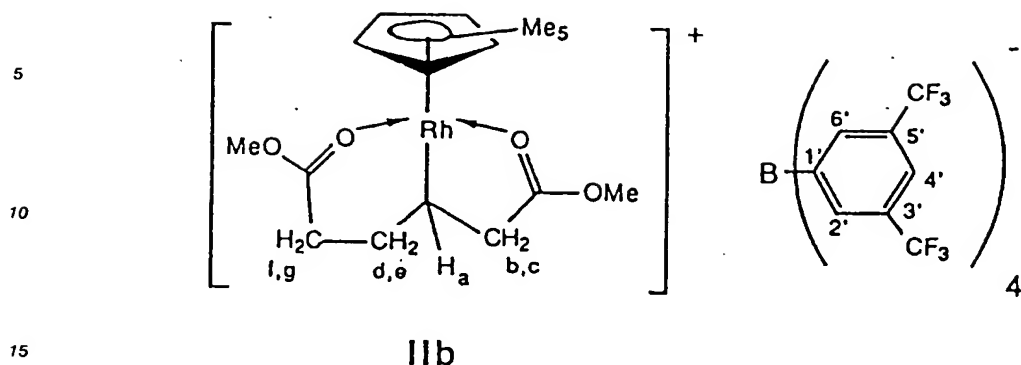
Method 1: HBPh₄⁺(Et₂O)₂ (218 mg, 0.215 mmol) in 3 mL CH₂Cl₂ was added at 0 °C to Cp*Rh(C₂H₄)₂ (49 mg, 0.167 mmol) in 7 mL CH₂Cl₂. After stirring 10 min, MeOC(O)CH=CHCH₂CH₂CO₂Me (200 μL) was added to the mixture. After stirring overnight at room temperature, the solution was evaporated to dryness. The residue was washed with hexane to eliminate the dimer. The two complexes (IIb) and (IIIb) were separated by successive recrystallizations in diethyl ether/hexane and isolated as orange crystals.

Method 2: HBPh₄⁺(Et₂O)₂ (171 mg, 0.169 mmol) in 3 mL CH₂Cl₂ was added at 0 °C to Cp*Rh(C₂H₄)₂ (39 mg, 0.133 mmol) in 7 mL CH₂Cl₂. After stirring 10 min, methyl acrylate (240 μL, 2.668 mmol) was added to the mixture. After stirring overnight at room temperature, the solution was evaporated to dryness. The residue was washed with hexane to eliminate excess dimer. The two complexes (IIb) and (IIIb) were separated by successive recrystallizations in diethyl ether/hexane and isolated as orange crystals.

NMR data for [Cp*RhCH(CH₂CO₂Me)(CH₂CH₂CO₂Me)]⁺[BPh₄⁻]⁻ (IIb):

¹H NMR (400 MHz, CD₂Cl₂, 23 °C): δ 7.72 (Ph⁺, 8H), 7.56 (Ph⁺, 4H), 3.93 (s, CO₂CH₃), 3.84 (s, CO₂CH₃), 3.35 (m, broad, Ha), 3.00 (dd, J_{Ha-Hb} or c = 9 Hz, J_{Hb-Hc} = 19 Hz, Hb or c), 2.68 (d, J_{Hb-Hc} = 19 Hz, Hc or b), 2.40 (ddd, J = 3, 7 and 19 Hz, Hf or g), 2.15 (ddd, J = 3, 9 and 19 Hz, Hg or f), 1.68 (m, Hd or e), 1.53 (s, C₅(CH₃)₅), 1.52 (m, He or d).

¹³C NMR (100 MHz, CD₂Cl₂, 23 °C): δ 190.4 (s, CO₂CH₃), 183.0 (s, CO₂CH₃), 162.1 (q, J_{C-B} = 50 Hz, C1'), 135.2 (d, J_{C-H} = 157.5 Hz, C2' and C6'), 129.3 (q, ²J_{C-F} = 32 Hz, C3' and C5'), 125.0 (q, J_{C-F} = 273 Hz, CF₃), 117.9 (dq, J_{C-H} = 163 Hz, ³J_{C-F} = 4 Hz, C4'), 94.6 (d, J_{C-Rh} = 8 Hz, C₅(CH₃)₅), 55.7 (q, J_{C-H} = 150 Hz, CO₂CH₃), 54.9 (q, J_{C-H} = 150 Hz, CO₂CH₃), 44.8 (t, J_{C-H} = 130 Hz, CH₂), 38.7 (dd, J_{C-Rh} = 23 Hz, J_{C-H} = 140 Hz, Rh-CH), 31.6 (t, J_{C-H} = 130 Hz, CH₂), 29.9 (t, J_{C-H} = 130, CH₂), 8.9 (q, J_{C-H} = 129 Hz, C₅-

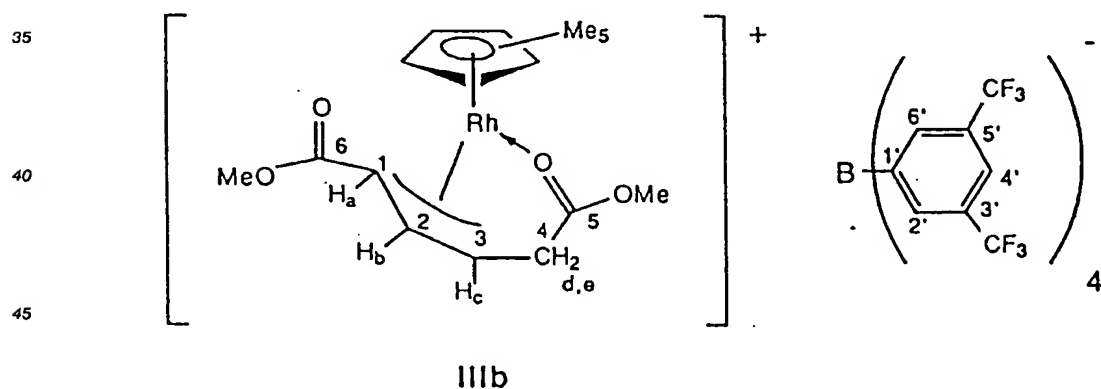
$(\text{CH}_3)_5$.

NMR data for $[\text{Cp}^*\text{Rh}(\eta^3\text{-MeOC(O)CH}_2\text{CHCHCHCO}_2\text{Me})]^+ [\text{BPh}_4]^-$ (IIb):

^1H NMR (400 MHz, CD_2Cl_2 , 23°C): δ 7.72 (Ph⁺, 8H), 7.56 (Ph⁺, 4H), 5.49 (ddd, $J_{\text{Hb-Hc}} = 11$ Hz, $J_{\text{Hc-Hd}} = 8$ Hz, $J_{\text{Rh-Hb}} = 2$ Hz, Hb), 4.70 (ddd, $J_{\text{Hb-Hc}} = 8$ Hz, $J_{\text{Hc-Hd}} = 7.5$ Hz, $J_{\text{Hc-He}} = 2$ Hz, Hc), 3.85 (s, CO_2CH_3), 3.82 (s, CO_2CH_3), 3.42 (dd, $J_{\text{Hd-Hc}} = 7.5$ Hz, $J_{\text{Hd-He}} = 21$ Hz, Hd), 3.11 (d, $J_{\text{Ha-Hb}} = 11$ Hz, Ha), 2.61 (dd, $J_{\text{He-Hd}} = 21$ Hz, $J_{\text{He-Hc}} = 2$ Hz, He), 1.70 (s, $\text{C}_5(\text{CH}_3)_5$).

$^{13}\text{C}\{^1\text{H}\}$ NMR (100 MHz, CD_2Cl_2 , 23°C): δ 186.8 (s, C5), 170.0 (s, C6), 162.1 (q, $J_{\text{C-B}} = 50$ Hz, C1'), 135.2 (s, C2' and C6'), 129.3 (q, $^2J_{\text{C-F}} = 32$ Hz, C3' and C5'), 125.0 (q, $J_{\text{C-F}} = 273$ Hz, CF₃), 117.9 (q, $^3J_{\text{C-F}} = 4$ Hz, C4'), 102.5 (d, $J_{\text{C-Rh}} = 5$ Hz, C2), 101.3 (d, $J_{\text{C-Rh}} = 7$ Hz, C5(CH₃)₅), 71.6 (d, $J_{\text{C-Rh}} = 9$ Hz, C3), 67.8 (d, $J_{\text{C-Rh}} = 10$ Hz, C1), 56.5 (s, OCH₃), 52.5 (s, OCH₃), 36.5 (s, C4), 8.9 (s, C₅(CH₃)₅).

^{13}C NMR (100 MHz, CD_2Cl_2 , 23°C): δ 186.8 (s, C5), 170.0 (s, C6), 162.1 (q, $J_{\text{C-B}} = 50$ Hz, C1'), 135.2 (d, $J_{\text{C-H}} = 157.5$ Hz, C2' and C6'), 129.3 (q, $^2J_{\text{C-F}} = 32$ Hz, C3' and C5'), 125.0 (q, $J_{\text{C-F}} = 273$ Hz, CF₃), 117.9 (dq, $J_{\text{C-H}} = 163$ Hz, $^3J_{\text{C-F}} = 4$ Hz, C4'), 102.5 (m, $J_{\text{C-H}} = 170$ Hz, C2), 101.3 (d, $J_{\text{C-Rh}} = 7$ Hz, C₅-(CH₃)₅), 71.6 (m, $J_{\text{C-H}} = 160$ Hz, C3), 67.8 (dt, $J_{\text{C-H}} = 161$ Hz, $^1J_{\text{C-Rh}} = ^2J_{\text{C-Hb}} = 10$ Hz, C1), 56.5 (q, $J_{\text{C-H}} = 151$ Hz, OCH₃), 52.5 (q, $J_{\text{C-H}} = 148$ Hz, OCH₃), 36.5 (t, $J_{\text{C-H}} = 130$ Hz, C4), 8.9 (q, $J_{\text{C-H}} = 129$ Hz, C₅-(CH₃)₅).



Claims

1. A process for preparing functionalized linear olefins which comprises reacting a first olefin, $\text{H}_2\text{C}=\text{CR}^1\text{R}^2$, with a second olefin, $\text{H}_2\text{C}=\text{CR}^3\text{R}^4$, in the presence of a cationic rhodium compound, $[\text{L}^1\text{RhL}^2\text{L}^3\text{R}]^+\text{X}^-$; wherein

R^1

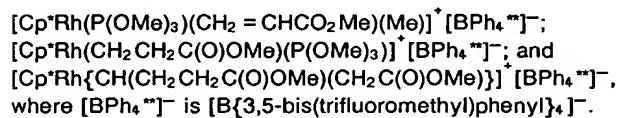
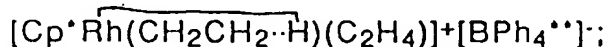
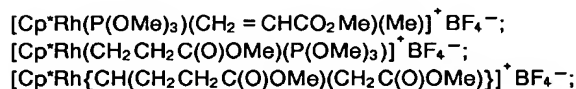
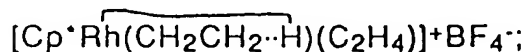
R^2

is selected from the group consisting of H and $\text{C}_1\text{-C}_{10}$ alkyl;

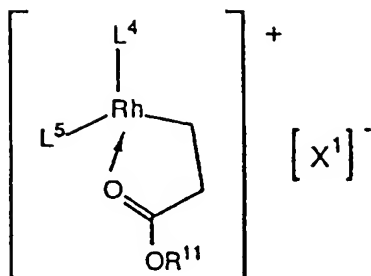
is selected from the group consisting of H, $\text{C}_1\text{-C}_{10}$ alkyl, phenyl, $\text{C}_7\text{-C}_{12}$ alkyl-substituted phenyl, $-\text{COOR}^5$, $-\text{C(O)NR}^6\text{R}^7$, and $-\text{C(O)H}$;

- R^3 is selected from the group consisting of H and C_1-C_{10} alkyl;
 R^4 is selected from the group consisting of $-COOR^8$, $-C(O)NR^9R^{10}$, and $-C(O)H$;
 R^5 and R^8 are independently selected from the group consisting of C_1-C_{10} alkyl;
 R^6 , R^7 , R^9 , and R^{10} are independently selected from the group consisting of H and C_1-C_{10} alkyl;
 L^1 is an anionic pentahapto ligand;
 L^2 and L^3 are neutral 2-electron donor ligands;
 R is selected from the group of H, C_1-C_{10} alkyl, C_6-C_{10} aryl, and C_7-C_{10} aralkyl ligands;
 X^- is a non-coordinating anion; and wherein two or three of L^2 , L^3 and R are optionally connected.

2. A process according to Claim 1 wherein said process is carried out between the temperatures of -100°C to 150°C .
3. A process according to Claim 2 wherein said first olefin is selected from the group consisting of ethylene, propylene, styrene, methyl acrylate, ethyl acrylate, acrolein, and N,N-dimethyl acrylamide and said second olefin is selected from the group consisting of methyl acrylate, ethyl acrylate, acrolein, and N,N-dimethyl acrylamide.
4. A process according to Claim 3 wherein L^1 is pentamethylcyclopentadienyl.
5. A process according to Claim 4 in which said first olefin is selected from the group consisting of ethylene, styrene, methyl acrylate and ethyl acrylate, and said second olefin is selected from the group consisting of methyl acrylate and ethyl acrylate.
6. A process according to Claim 5 in which said cationic rhodium compound is selected from the group consisting of:



7. A process according to Claim 6 in which the partial pressure of hydrogen is 0.1 to 10 atm.
8. A process according to Claim 7 in which said first olefin is methyl acrylate and said second olefin is methyl acrylate.
9. A compound of the formula



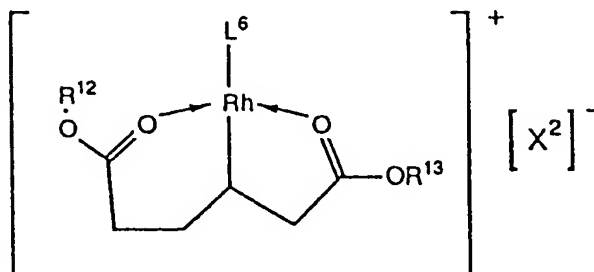
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11. A compound according to Claim 10 wherein L⁵ is trimethylphosphite and R¹¹ is methyl.

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